### CLAIMS

- 1. A method for inducing immunosuppressive cells, which comprises culturing human cells with the use of a culture device having an affinity for protein.
- 2. The inducing method of Claim 1, wherein the culture device is previously coated with one or more cytokines or antibodies against surface antigens.

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3. The inducing method of Claim 1, wherein the culture device is previously coated with two or more antibodies against surface antigens, each of said antibodies recognizing different epitope.

4. The inducing method of Claim 1, wherein human cells and one or more cytokines or antibodies against surface antigens are mixed.

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5. The inducing method of Claim 1, wherein human cells and two or more antibodies against surface antigens, each of said antibodies recognizing different epitope, are mixed.

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6. The inducing method of Claim 2, 3, 4 or 5, wherein the antibody against surface antigens is an anti-CD2 antibody or an anti-CD3 antibody.

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8. The inducing method of Claim 2, 3, 4, 5 or 6, wherein the antibody against surface antigens is an anti-CD2 antibody, said anti-CD2 antibody binding to a site of CD2 other than a site of CD2 which participates in the binding of LFA-3 to CD2.

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9. The inducing method of Claim 3 or 5, wherein two or more antibodies against surface antigens are anti-CD2 antibodies comprising a combination of one or more antibodies which bind to a site of CD2 that participates in the binding of LFA-3 to CD2 and one or more antibodies which bind to a site of CD2 other than the site of CD2 which participates in the binding of LFA-3 to CD2.

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10. The inducing method of Claim 2, 3, 4, 5, 6, 7 or 8, wherein one antibody against surface antigens is the anti-CD2 antibody TS2/18.

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fragment of the anti-CD2 antibody TS2/18.

wherein one antibody against surface antigens is the F(ab)2

11. The inducing method of Claim 2, 3, 4, 5, 6, 7 or 8,

12. The inducing method of Claim 2 or 4, wherein the cytokine is IL-2 or GM-CSF.

- 13. The inducing method of Claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11 or 12, wherein 0.5 to 10 % by volume of serum based on a culture medium is mixed in the culture.
- 14. The inducing method of Claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11 or 12, wherein no serum is mixed with the culture medium in the culture.
- 15. The inducing method of Claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10 10, 11, 12, 13 or 14, wherein a term of the culture ranges between 1 and 7 days.

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- 16. The inducing method of Claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15, wherein the culture device is made of material having an affinity for protein.
- 17. The inducing method of Claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15 or 16, wherein the culture device is made of plastic material.
- 18. The inducing method of Claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15 or 16, wherein the culture device is made of glass material.
- 19. The inducing method of Claim 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17 or 18, wherein the culture device is a closed plane plate vessel or a closed vessel charged with

spherical fine-particles.

- 20. A culture device having an affinity for protein, which is used for inducing immunosuppressive cells by culturing human cells.
- 21. The culture device of Claim 20, wherein the device is previously coated with one or more cytokines or antibodies against surface antigens.

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22. The culture device of Claim 20, wherein the device is previously coated with two or more antibodies against surface antigens, each of said antibodies recognizing different epitope.

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- 23. The culture device of Claim 21 or 22, wherein the antibody against surface antigens is anti-CD2 antibody or anti-CD3 antibody.
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- 24. The culture device of Claim 22 or 23, wherein one or more antibodies that recognize different epitope are the anti-CD2 antibody TS2/18.

## **IN THE CLAIMS:**

#### Please cancel claims 1-19

#### Please amend the claims as follows:

- 20. A culture device having an affinity for protein, which is used for inducing immunosuppressive cells by culturing human cells <u>in said device</u>.
- 21. The culture device of Claim 20, wherein <u>prior to said culturing</u> the device is [previously] coated with one or more cytokines <u>and/or one or more</u> antibodies against <u>cell</u> surface antigens.
- 22. The culture device of Claim 20, wherein <u>prior to said culturing</u> the device is [previously] coated with two or more antibodies against <u>cell</u> surface antigens, each of said antibodies recognizing <u>a</u> different epitope.
- 23. The culture device of Claim 21 [or 22], wherein <u>said antibodies</u> [the antibody] against <u>cell</u> surface antigens <u>are</u> [is] anti-CD2 <u>antibodies</u> [antibody] or anti-CD3 <u>antibodies</u> [antibody].
- 24. The culture device of Claim 23 or 25, wherein [one or more antibodies that recognize different epitope are] the anti-CD2 antibody is TS2/18 produced from hybridoma HB195 (ATCC Accession number HB-195).

#### Please add new claim 25:

25. The culture device of Claim 22, wherein said antibodies against cell surface antigens are anti-CD2 antibodies or anti-CD3 antibodies.

# PRELIMINARY AMENDMENT Of Parent U.S. Appln. No. 09/254,170

## **REMARKS**

This response follows an Office Action of September 26, 2000 withdrawing claims 20-

25. Claims 1-19 are all the claims pending in the parent application.

Entry and consideration of this Amendment is respectfully requested.

Respectfully submitted,

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